## SOLMEET DENTI DOCTOR - calcium carbonate paste, dentifrice Solbin Co., Ltd

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

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## **Drug Facts**

Active Ingredient: Calcium Carbonate, Dibasic Calcium Phosphate Hydrate, Aminocarproic Acid, Aluminium Chlorohydroxy Allantoinate

Inactive ingredients: calcium carbonate, dibasic calcium phosphate hydrate, amonicaproic acid, aluminium chlorohydroxy allantoinate, glycerine, D-sorbitol, polyethylene glycol-1500, carboxymethylcellulose sodium, sodium lauryl sulfate, methylparaben, xylitol, L-menthol, peppermint oil, yellow no.5, blue no.1, angelica dahurica root extract, phellodendron bark extract, rosin, purified water

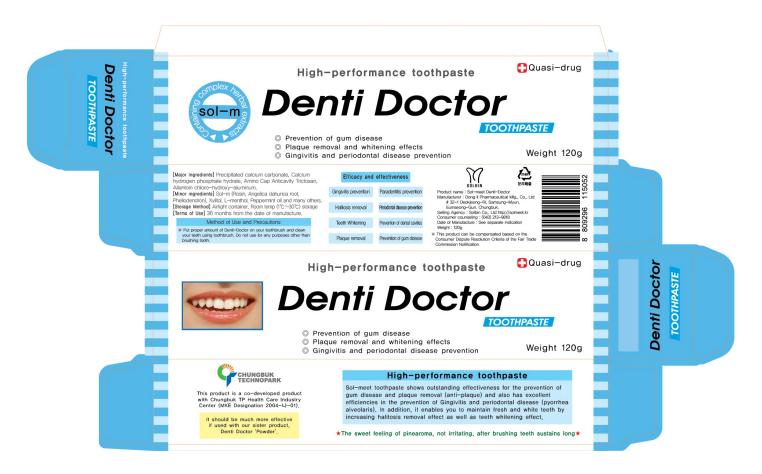
for dental care

keep out of reach of the children

use when is needed

- do not swallow when using this product
- if more than used for rinsing is accidentally swallowed, get medical helps or contact a poison control center right away

apply proper amount on your toothbrush



calcium carbonate paste, dentifrice

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:42352-8001
Route of Administration	DENTAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
CALCIUM CARBONATE (UNII: H0 G9 379 FGK) (CARBONATE ION - UNII:7UJQ5OPE7D)	CALCIUM CARBONATE	15.85 g in 100 g	
<b>DIBASIC CALCIUM PHO SPHATE DIHYDRATE</b> (UNII: O7TSZ97GEP) (PHOSPHATE ION - UNII:NK08V8K8HR)	DIBASIC CALCIUM PHOSPHATE DIHYDRATE	0.6 g in 100 g	
<b>AMINO CAPRO IC ACID</b> (UNII: U6F3787206) (AMINO CAPRO IC ACID - UNII: U6F3787206)	AMINOCAPROIC ACID	0.05 g in 100 g	
ALCLOXA (UNII: 18B8O9DQA2) (ALCLOXA - UNII:18B8O9DQA2)	ALCLOXA	0.05 g in 100 g	

Inactive Ingredients		
Ingredient Name	Strength	
SODIUM CHLORIDE (UNII: 451W47IQ8X)		
GLYCERIN (UNII: PDC6A3C0OX)		
SORBITOL (UNII: 506T60A25R)		
POLYETHYLENE GLYCOL 1500 (UNII: 1212Z7S33A)		
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311)		
SODIUM LAURYL SULFATE (UNII: 368GB5141J)		
METHYLPARABEN (UNII: A2I8C7HI9T)		
XYLITOL (UNII: VCQ006KQ1E)		
LEVOMENTHOL (UNII: BZ1R15MTK7)		
PEPPERMINT (UNII: V95R5KMY2B)		
FD&C YELLOW NO. 5 (UNII: 1753WB2F1M)		
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)		
ANGELICA DAHURICA ROOT (UNII: 1V63N2S972)		
PHELLODENDRON AMURENSE BARK (UNII: PBG27B754G)		
<b>ROSIN</b> (UNII: 88S87KL877)		
WATER (UNII: 059QF0KO0R)		

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:42352-8001-1	120 g in 1 TUBE		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
unapproved drug other		12/15/2010		

## **Labeler -** Solbin Co., Ltd (631099371)

## Registrant - Solbin Co., Ltd (631099371)

Establishment				
Name	Address	ID/FEI	Business Operations	
Solbin Co., Ltd		631099371	manufacture	

Revised: 12/2010 Solbin Co., Ltd